AMENDMENT UNDER 37 C.F.R. § 1.111

Application No.: 10/583,849

AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the

application:

LISTING OF CLAIMS:

1. (currently amended): A process for producing a fibre composition comprising a

lignocellulosic fibre material containing phenolic or similar structural groups capable of

being oxidized, and a synthetic, electrically conductive polymer formed by polymerized

monomers, according to which process the monomers are polymerized in the presence of the

lignocellulosic fibre material to form a composition in which the polymer is bound to the

fibres, characterized by

a) oxidizing the phenolic groups or the groups having a similar structure to

provide an oxidized fibre material,

b) contacting the oxidized fibre material with a bifunctional substance to provide

a modified lignocellulosic fibre material capable of binding monomers of the

conductive polymer, and

c) contacting the modified lignocellulosic fibre material with monomers of the

conductive polymer under conditions conducive to polymerization to produce

polymer chains of the synthetic, electrically conductive polymer, which are

grafted to the surface of the lignocellulosic fibre material,

wherein the bifunctional substance has at least two functional groups, where the first

functional group participates in the binding of the modifying compound to the lignocellulosic

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fibre material and the second functional group forms a primer for binding to the polymeric

material.

2. (original): The process according to claim 1, wherein the oxidized fibre material is contacted

with the bifunctional monomers of the synthetic, electrically conductive polymer in order to

bind the monomers to the surface of the oxidized lignocellulosic fibre material, to provide a

modified lignocellulosic fibre material having monomers bound to its surface, and the

modified lignocellulosic fibre material is contacted with the monomers to produce polymer

chains of the synthetic, electrically conductive polymer, which are grafted to the surface of

the lignocellulosic fibre material.

3. (original): The process according to claim 1, wherein the modifying agent is activated with

an oxidizing agent.

4. (currently amended): The process according to claim 1, wherein the lignocellulosic fibrous

matrix material is reacted with an oxidizing agent in the presence of a substance capable of

catalyzing the oxidation of phenolic groups or groups having a similar structure by said

oxidizing agent.

5. (previously presented): The process according to claim 1, wherein the reaction of step (a) is

carried out in an aqueous phase at a consistency of about 1 to 95 % by weight of the fibre

material.

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6. (currently amended): The process according to claim 1, wherein the polymer is selected from the group consisting of polyaniline, polypyrrole, polythiophene and polyacetylene and derivatives thereof.

7. (cancelled)

- 8. (currently amended): The process according to claim 71, wherein the modifying agent comprises at least one phenolic hydroxyl or similar structural group as a first functional group.
- 9. (currently amended): The process according to claim 71, wherein the second functional group is selected from the group of hydroxy, carboxy, anhydride, aldehyde, ketone, amine, amide, imine, imidine and derivatives and salts thereof.
- 10. (currently amended): The process according to claim $7\underline{1}$, wherein the modifying agent comprises a plurality of second functional groups.
- 11. (previously presented): The process according to claim 1, wherein the bifunctional substance and the monomer are different.

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12. (previously presented): The process according to claim 1, wherein the bifunctional substance

and the monomer are the same.

13. (previously presented): The process according to claim 1, wherein the fibres are selected

from lignocellulosic fibres produced by mechanical, chemimechanical or chemical pulping.

14. (previously presented): The process according to claim 4, wherein the substance capable of

catalyzing the oxidation of phenolic or similar structural groups to provide an oxidized fibre

material is an enzyme.

15. (previously presented): The process according to claim 1, wherein steps (a) to (c) are carried

out simultaneously by forming in an aqueous medium a mixture of lignocellulosic fibres and

the monomer, oxidizing phenolic or similar structural groups on the lignocellulosic fibres

while binding the monomers to the oxidized phenolic or similar structural groups.

16. (currently amended): The process according to claim 1514, wherein the enzyme is added to

the aqueous medium in order to oxidized the phenolic or similar structural groups.

17. (previously presented): The process according to claim 14, wherein the enzyme capable of

catalyzing the oxidation of phenolic groups is selected from the group of peroxidases and

oxidases.

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18. (currently amended): The process according to claim 1714, wherein the enzyme is selected the group of laccases (EC 1.10.3.2), catechol oxidases (EC 1.10.3.1), tyrosinases (EC 1.14.18.1), bilirubin oxidases (EC 1.3.3.5), horseradish peroxidase (EC 1.11.1.7).

- 19. (currently amended): The process according to claim 17, wherein the enzyme dosage is from about 1 to 100,000 nkat/g, preferably 10 500 nkat/g, and it is employed in an amount of 0.0001 to 10 mg protein/g of dry matter.
- 20. (currently amended): The process according to claim 19, wherein the enzyme treatment is carried out at a temperature of 5-100 °C, preferably 10-85 °C and most preferably at 20-80 °C and pH 3-12.
- 21. (currently amended): The process according to claim 3, wherein the oxidizing agent is selected from the group of oxygen and oxygen-containing gases, such as selected from the group consisting of air and hydrogen peroxide.
- 22. (original): The process according to claim 21, wherein oxygen or oxygen-containing gas or hydrogen peroxide is introduced into the aqueous slurry during the reaction.
- 23. (previously presented): The process according to claim 1, wherein a chemical oxidizing agent is used.

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24. (currently amended): The process according to claim 22, wherein the chemical oxidizing agent is hydrogen peroxide, Fenton reagent, potassium permanganate, ozone and chloride dioxide or an inorganic transition metal salt, ammoniumperoxy sulphate.

- 25. (currently amended) The process according to claim 1, wherein <u>said oxidizing step comprises</u>

 <u>radiating the lignocellulosic fibre matrix employing</u> radical forming radiation capable of

 catalyzing the oxidation of phenolic or similar structural groups <u>is used</u> to provide an

 oxidized fibre material.
- 26. (previously presented): The process according to claim 1, wherein the reaction steps are carried out sequentially or simultaneously.
- 27. (new): The process according to claim 19, wherein the enzyme dosage is from 10-500 nkat/g.
- 28. (new): The process according to claim 20, wherein the enzyme treatment is carried out at a temperature of 10-85°C.
- 29. (new): The process according to claim 20, wherein the enzyme treatment is carried out at a temperature of 20-80°C.

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30. (new): The process according to claim 6, wherein the derivatives are selected from alkyl derivatives, aryl derivatives, chlorine substituted derivatives, or bromine substituted

derivatives.